## **AMENDMENTS TO THE CLAIMS**

This Listing of Claims will replace all prior versions, and listings, of claims in the application.

## **Listing Of Claims:**

Claim 1 (Currently Amended): <u>A process Process</u> for the preparation of enantiomerically pure (S)- or (R) -4-halo-3-hydroxybutyrates of formula:

$$R^1$$
  $OH$   $O$   $OR^2$   $(S)$ -I or  $R^1$   $OR^2$   $(R)$ -I,

wherein  $R^1$  is  $CH_2X$ ,  $CHX_2$  or  $CX_3$  and X independently represents CI and/or Br and wherein  $R^2$  is  $C_{1-6}$ -alkyl,  $C_{3-6}$ -cycloalkyl, aryl or aralkyl, each aryl or aralkyl being optionally further substituted with one or more  $C_{1-4}$ -alkyl groups and/or halogen atoms, which process comprises the asymmetric hydrogenation of 4-halo-3-oxobutyates of formula:

wherein R1, R2 and X are as defined above

in the presence of a catalyst of a ruthenium complex comprising a chiral ligand of formula:

$$\begin{array}{c} F \\ O \\ \hline \\ F \\ O \\ \hline \end{array}$$
 PPh<sub>2</sub> III.

said resultant (S)- or (R)-4-halo-hydroxybutyrate has an enantiomeric purity in the range of an ee of 93.2 percent to an ee of 98.1 percent.

Claim 2 (Previously Presented): The process of claim 1, wherein the ruthenium complex comprising a ligand of formula III comprises at least one diene, alkene or arene or polar solvent molecule as stabilizing ligand.

Claim 3 (Previously Presented): The process of claim 1, wherein the ruthenium complex comprising a ligand of formula III comprises at least one molecule of 1,5-cyclooctadiene or p-cymene as stabilizing ligand.

Claim 4 (Previously Presented): The process of claim 1, wherein the hydrogenation is carried out in a solution comprising a polar solvent selected from the group consisting of C<sub>1-4</sub>-alcohols, dimethylsulfoxide, demethylformamide, acetonitrile and mixtures thereof, wherein the solvent optionally contains further solvent additives.

Claim 5 (Previously Presented): The process of claim 1, wherein the counterion of the ruthenium complex is selected from the group consisting of  $Cl^{-}$ ,  $Br^{-}$ ,  $l^{-}$ ,  $BF_{4}^{-}$ ,  $AsF_{6}^{-}$ ,  $SbF_{6}^{-}$ ,  $PF_{6}^{-}$ ,  $ClO_{4}^{-}$  and  $OTf^{-}$ .

Claim 6 (Previously Presented): The process of claim 1, wherein the ruthenium complex is prepared by mixing the complex of formula [Ru<sub>2</sub>Cl<sub>4</sub>(cym)<sub>2</sub>] with the Fluoxphos ligand in a polar solvent.

Claim 7 (Currently Amended): The process of claim 1, wherein the hydrogen pressure during the reaction is in the range of 1 to 60 and <del>preferably in the range of 2 to 35 bar.</del>

Claim 8 (Previously Presented): The process of claim 2, wherein the ruthenium complex comprising a ligand of formula III comprises at least one molecule of 1,5-cyclooctadiene or *p*-cymene as stabilizing ligand.

Claim 9 (Currently Amended): The process of claim 2, wherein the hydrogenation is carried out in a solution comprising a polar solvent selected from the group consisting of C<sub>1-4</sub>-alcohols, dimethylsulfoxide, dimethylformamide, acetonitrile and mixtures thereof, wherein the <u>polar</u> solvent optionally contains further solvent additives.

Claim 10 (Currently Amended): The process of claim 3, wherein the hydrogenation is carried out in a solution comprising a polar solvent selected from the group consisting of C<sub>1-4</sub>-alcohols, dimethylsulfoxide, dimethylformamide, acetonitrile and mixtures thereof, wherein the polar solvent optionally contains further solvent additives.

Claim 11 (Previously Presented): The process of claim 2, wherein the counterion of the ruthenium complex is selected from the group consisting of  $Cl^-$ ,  $Br^-$ ,  $l^-$ ,  $BF_4^-$ ,  $AsF_6^-$ ,  $SbF_6^-$ ,  $PF_6^-$ ,  $ClO_4^-$  and  $OTf^-$ .

Claim 12 (Previously Presented): The process of claim 8, wherein the counterion of the ruthenium complex is selected from the group consisting of Cl $^-$ , Br $^-$ , I $^-$ , BF $_4$  $^-$ , AsF $_6$  $^-$ , SbF $_6$  $^-$ , PF $_6$  $^-$ , ClO $_4$  $^-$  and OTf $^-$ .

Claim 13 (Previously Presented): The process of claim 10, wherein the counterion of ruthenium complex is selected from the group consisting of Cl $^-$ , Br $^-$ , I $^-$ , BF $_4$  $^-$ , AsF $_6$  $^-$ , SbF $_6$  $^-$ , PF $_6$  $^-$ , ClO $_4$  $^-$  and OTf $^-$ .

Claim 14 (Previously Presented): The process of claim 9, wherein the counterion of ruthenium complex is selected from the group consisting of Cl $^-$ , Br $^-$ , l $^-$ , BF $_4$  $^-$ , AsF $_6$  $^-$ , SbF $_6$  $^-$ , PF $_6$  $^-$ , ClO $_4$  $^-$  and OTf $^-$ .

Claim 15 (Previously Presented): The process of claim 2, wherein the ruthenium complex is prepared by mixing the complex of formula  $[Ru_2Cl_4(cym)_2]$  with the Fluoxphos ligand in a polar solvent.

Claim 16 (Previously Presented): The process of claim 8, wherein the ruthenium complex is prepared by mixing the complex of formula [Ru<sub>2</sub>Cl<sub>4</sub>(cym)<sub>2</sub>] with the Fluoxphos ligand in a polar solvent.

Claim 17 (Previously Presented): The process of claim 9, wherein the ruthenium complex is prepared by mixing the complex of formula [Ru<sub>2</sub>Cl<sub>4</sub>(cym)<sub>2</sub>] with the Fluoxphos ligand in a polar solvent.

Claim 18 (Previously Presented): The process of claim 14, wherein the ruthenium complex is prepared by mixing the complex of formula [Ru<sub>2</sub>Cl<sub>4</sub>(cym)<sub>2</sub>] with the Fluoxphos ligand in a polar solvent.

Claim 19 ((Currently Amended): The process of claim 2, wherein the hydrogen pressure during the reaction is in the range of 1 to 60 bar and preferably in the range of 2 to 35 bar.

Claim 20 (Currently Amended): The process of claim 8, wherein the hydrogen pressure during the reaction is in the range of 1 to 60 bar and preferably in the range of 2 to 35 bar.

Claim 21 (Currently Amended): The process of claim 9, wherein the hydrogen pressure during the reaction is in the range of 1 to 60 bar and preferably in the range of 2 to 35.

Claim 22 (Currently Amended): The process of claim 14, wherein the hydrogen pressure during the reaction is in the range of 1 to 60 bar and preferably in the range of 2 to 35 bar.

Claim 23 (Currently Amended): The process of claim <u>1</u> 48, wherein the hydrogen pressure during the reaction is in the range of <u>1 to 60 bar and</u> preferably in the range of 2 to 35 bar.